

**Amendments to the Claims:**

This listing of claims will replace all prior versions, and listings, of claims in the application:

**Listing of Claims:**

1-88. (Canceled)

89. (Original) A method for determining whether a candidate molecule affects a body weight disorder associated with an organism, comprising:

(a) contacting a cell from said organism with, or recombinantly expressing within the cell from said organism, said candidate molecule;

(b) determining whether the RNA expression or protein expression in said cell of at least one open reading frame is changed in step (a) relative to the expression of said open reading frame in the absence of the candidate molecule, each said open reading frame being regulated by a promoter native to a nucleic acid sequence selected from the group consisting of SEQ ID NO: 5, SEQ ID NO: 6, SEQ ID NO: 7, SEQ ID NO: 8, SEQ ID NO: 9, SEQ ID NO: 11, SEQ ID NO: 12, SEQ ID NO: 14, SEQ ID NO: 16, SEQ ID NO: 18, SEQ ID NO: 20, SEQ ID NO: 21, SEQ ID NO: 23 and homologs of each of the foregoing; and

(c) determining that the candidate molecule affects a body weight disorder associated with said organism when the RNA expression or protein expression of said at least one open reading frame is changed, or

determining that the candidate molecule does not affect a body weight disorder associated with said organism when the RNA expression or protein expression of said at least one open reading frame is unchanged.

90-107. (Canceled)

108. (Original) A method for determining whether a first trait  $T_1$  is causal for a second trait  $T_2$  in a plurality of organisms of a species, the method comprising:

(A) identifying one or more loci in the genome of said species, wherein each locus  $Q$  of said one or more loci is a site of colocalization for (i) a respective quantitative trait locus

(QTL<sub>1</sub>) that is genetically linked to a variation in the first trait T<sub>1</sub> across the plurality of organisms and (ii) a respective quantitative trait locus (QTL<sub>2</sub>) that is genetically linked to a variation in the second trait T<sub>2</sub> across said plurality of organisms; and

(B) testing, for each respective locus  $Q$  of said one or more loci, whether (i) a genetic variation  $Q^*$  of said respective locus  $Q$  across said plurality of organisms and (ii) said variation in said second trait T<sub>2</sub> across said plurality of organisms are correlated conditional on said variation in said first trait T<sub>1</sub> across said plurality of organisms,

wherein, when the genetic variation of (i) one or more loci  $Q$  tested in (B), and (ii) said variation in said second trait T<sub>2</sub> across said plurality of organisms are correlated conditional on said variation in said first trait T<sub>1</sub> across said plurality of organisms, said first trait T<sub>1</sub> is determined to be causal for said second trait T<sub>2</sub>.

109. (Original) The method of claim 108, the method further comprising, prior to said identifying, a step of determining a respective QTL<sub>1</sub> at a locus  $Q$  of said one or more loci using a first quantitative trait locus (QTL) analysis, wherein said first QTL analysis uses a plurality of quantitative measurements of said first trait, and wherein each quantitative measurement in said plurality of quantitative measurements of said first trait is associated with an organism in said plurality of organisms.

110. (Original) The method of claim 109, the method further comprising a step of determining a respective QTL<sub>2</sub> at said locus  $Q$  using a second QTL analysis, wherein said second QTL analysis uses a plurality of quantitative measurements of said second trait, and wherein each quantitative measurement in said plurality of quantitative measurements of said second trait is associated with an organism in said plurality of organisms.

111. (Original) The method of claim 108, wherein said respective QTL<sub>1</sub> and said respective QTL<sub>2</sub> are deemed to be colocalized at a locus  $Q$  of said one or more loci when said respective QTL<sub>1</sub> and said respective QTL<sub>2</sub> are within 3 cM of the locus  $Q$ .

112. (Original) The method of claim 108, wherein said respective QTL<sub>1</sub> and said respective QTL<sub>2</sub> are deemed to be colocalized at a locus  $Q$  of said one or more loci when said respective QTL<sub>1</sub> and said respective QTL<sub>2</sub> are within 1 cM of the locus  $Q$ .

113. (Original) The method of claim 108 wherein said plurality of organisms is derived from a predetermined set of individuals.

114. (Original) The method of claim 108 wherein said plurality of organisms is derived from a predetermined set of strains.

115. (Original) The method of claim 114 wherein said set of strains is between 2 strains and 100 strains.

116. (Original) The method of claim 114 wherein said set of strains is between 5 strains and 500 strains.

117. (Original) The method of claim 114 wherein said set of strains is more than five strains.

118. (Original) The method of claim 114 wherein said set of strains is less than 1000 strains.

119. (Original) The method of claim 114 wherein said set of strains is diverse with respect to a complex phenotype associated with human disease.

120. (Original) The method of claim 114 wherein said set of strains is between 2 strains and 10 strains that, collectively, are diverse with respect to a complex phenotype associated with a human disease.

121. (Original) The method of claim 120 wherein said human disease is obesity, diabetes, atherosclerosis, metabolic syndrome, depression, anxiety, osteoporosis, bone development, asthma, or chronic obstructive pulmonary disease.

122. (Original) The method of claim 108 wherein said plurality of organisms is derived from crossing a predetermined set of strains.

123. (Original) The method of claim 122 wherein said plurality of organisms is an F<sub>2</sub> intercross, a backcross, or an F<sub>2</sub> random mating.

124. (Original) The method of claim 108 wherein the plurality of organisms is more than 1,000 organisms.

125. (Original) The method of claim 108 wherein the plurality of organism is between 100 organisms and 100,000 organisms.

126. (Original) The method of claim 108 wherein the plurality of organisms is less than 500,000 organisms.

127. (Original) The method of claim 108 wherein the plurality of organisms is between 5,000 and 25,000 organisms.

128. (Original) The method of claim 109, wherein

said first trait is abundance levels of a first cellular constituent and each quantitative measurement of said first trait is an abundance level of said first cellular constituent in an organism in said plurality of organisms; and

said second trait is abundance levels of a second cellular constituent and each quantitative measurement of said second trait is an abundance level of said second cellular constituent in an organism in said plurality of organisms.

129. (Original) The method of claim 128 wherein each said abundance level of said first cellular constituent is normalized and each said abundance level of said second cellular constituent is normalized

130. (Original) The method of claim 128 wherein

each said abundance level of said first cellular constituent is determined by measuring an amount of said first cellular constituent in one or more cells from an organism in said plurality of organisms; and

each said abundance level of said second cellular constituent is determined by measuring an amount of said second cellular constituent in one or more cells from an organism in said plurality of organisms.

131. (Original) The method of claim 128, wherein

each said amount of said first cellular constituent comprises an abundance of a first RNA in said one or more cells of said organism in said plurality of organisms; and

each said amount of said second cellular constituent comprises an abundance of a second RNA in said one or more cells of said organism in said plurality of organisms.

132. (Original) The method of claim 131, wherein

said abundance of said first RNA is measured by contacting a gene transcript array with said first RNA from said one or more cells of said organism, or with nucleic acid derived from said first RNA, wherein said gene transcript array comprises a positionally addressable surface with attached nucleic acids or nucleic acid mimics, wherein said nucleic acids or nucleic acid mimics are capable of hybridizing with said first RNA, or with nucleic acid derived from said first RNA; and

said abundance of said second RNA is measured by contacting a gene transcript array with said second RNA from said one or more cells of said organism, or with nucleic acid derived from said second RNA, wherein said gene transcript array comprises a positionally addressable surface with attached nucleic acids or nucleic acid mimics, wherein said nucleic acids or nucleic acid mimics are capable of hybridizing with said second RNA, or with nucleic acid derived from said second RNA.

133. (Original) The method of claim 109, wherein said first QTL analysis comprises:

(i) testing for linkage between (a) the genotype of said plurality of organisms at a position in the genome of said species and (b) said plurality of quantitative measurements of said first trait;

(ii) advancing the position in said genome by an amount; and

(iii) repeating steps (i) and (ii) until all or a portion of the genome of said species has been tested.

134. (Original) The method of claim 110, wherein said second QTL analysis comprises:

(i) testing for linkage between (a) the genotype of said plurality of organisms at a position in the genome of said species and (b) said plurality of quantitative measurements of said second trait;

(ii) advancing the position in said genome by an amount; and

(iii) repeating steps (i) and (ii) until all or a portion of the genome of said species has been tested.

135-136. (Cancelled)

137. (Original) The method of claim 133 or 134, wherein said testing comprises performing linkage analysis or association analysis.

138. (Original) The method of claim 137, wherein said linkage analysis or association analysis generates a statistical score for said position in the genome of said species.

139. (Original) The method of claim 138, wherein said testing is linkage analysis and said statistical score is a logarithm of the odds (lod) score.

140-141. (Cancelled)

142. (Original) The method of claim 109, wherein said respective  $QTL_1$  is represented by a lod score that is greater than 4.0.

143. (Original) The method of claim 110, wherein said respective  $QTL_2$  is represented by a lod score that is greater than 4.0.

144. (Original) The method of claim 109 wherein each quantitative measurement in said plurality of quantitative measurements of said first trait is

an amount or a concentration of a first cellular constituent in one or more tissues of an organism in said plurality of organisms,



a cellular constituent activity level of said first cellular constituent in one or more tissues of an organism in said plurality of organisms, or

a state of cellular constituent modification of said first cellular constituent in one or more tissues of an organism in said plurality of organisms.

145. (Original) The method of claim 110 wherein each quantitative measurement in said plurality of quantitative measurements of said second trait is

an amount or a concentration of a second cellular constituent in one or more tissues of an organism in said plurality of organisms,

a cellular constituent activity level of said second cellular constituent in one or more tissues of an organism in said plurality of organisms, or

a state of cellular constituent modification of said second cellular constituent in one or more tissues of an organism in said plurality of organisms.

146. (Original) The method of claim 108, wherein said plurality of organisms is human.

147. (Original) The method of claim 109, wherein said respective  $QTL_1$  and said respective  $QTL_2$  are deemed to colocalize at a locus  $Q$  of said one or more loci when said respective  $QTL_1$  and said respective  $QTL_2$  are within 40 cM of the locus  $Q$ .

148. (Original) The method of claim 109, wherein said respective  $QTL_1$  and said respective  $QTL_2$  are deemed to colocalize at a locus  $Q$  of said one or more loci when said respective  $QTL_1$  and said respective  $QTL_2$  are within 10 cM of the locus  $Q$ .

149. (Original) The method of claim 108 wherein said one or more loci consist of at least two loci.

150. (Original) The method of claim 108, wherein said respective  $QTL_1$  and said respective  $QTL_2$  colocalize at a locus  $Q$  of said one or more loci when said respective  $QTL_1$  and said respective  $QTL_2$  satisfy a pleiotropy test and wherein failure of the pleiotropy test indicates that (i) the respective  $QTL_1$  and the respective  $QTL_2$  are two closely linked QTL, (ii) step (B)

is not performed, and (iii) said first trait  $T_1$  is not determined to be causal for said second trait  $T_2$ .

151. (Original) The method of claim 150 wherein said pleiotropy test comprises comparing a model for a null hypothesis, indicating that said respective  $QTL_1$  and said respective  $QTL_2$  colocalize as a QTL, to a model for an alternative hypothesis, indicating that said  $QTL_1$  and said respective  $QTL_2$  are two closely linked QTL.

152. (Original) The method of claim 151 wherein said model for said null hypothesis is:

$$\begin{pmatrix} y_1 \\ y_2 \end{pmatrix} = \begin{pmatrix} \mu_1 \\ \mu_2 \end{pmatrix} + \begin{pmatrix} \beta_1 \\ \beta_2 \end{pmatrix} N + \begin{pmatrix} \varepsilon_1 \\ \varepsilon_2 \end{pmatrix}$$

wherein

$N$  is a categorical random variable indicating the genotype at locus  $Q$  across said plurality of organisms;

$\begin{pmatrix} \varepsilon_1 \\ \varepsilon_2 \end{pmatrix}$  is distributed as a bivariate normal random variable with mean  $\begin{pmatrix} 0 \\ 0 \end{pmatrix}$  and

covariance matrix  $\begin{pmatrix} \sigma_1^2 & \sigma_1\sigma_2 \\ \sigma_2\sigma_1 & \sigma_2^2 \end{pmatrix}$ ; and

$\mu_i$  and  $\beta_i$  are model parameters.

153. (Original) The method of claim 151 wherein said model for said alternative hypothesis is:

$$\begin{pmatrix} y_1 \\ y_2 \end{pmatrix} = \begin{pmatrix} \mu_1 \\ \mu_2 \end{pmatrix} + \begin{pmatrix} \beta_1 & \beta_2 \\ \beta_3 & \beta_4 \end{pmatrix} \begin{pmatrix} N_1 \\ N_2 \end{pmatrix} + \begin{pmatrix} \varepsilon_1 \\ \varepsilon_2 \end{pmatrix}$$

wherein

$N_1$  and  $N_2$  are categorical random variables indicating the genotype at locus  $Q$  across said plurality of organisms;

$\begin{pmatrix} \varepsilon_1 \\ \varepsilon_2 \end{pmatrix}$  is distributed as a bivariate normal random variable with mean  $\begin{pmatrix} 0 \\ 0 \end{pmatrix}$  and

covariance matrix  $\begin{pmatrix} \sigma_1^2 & \sigma_1\sigma_2 \\ \sigma_2\sigma_1 & \sigma_2^2 \end{pmatrix}$ ; and



$\mu_i$  and  $\beta_i$  are model parameters.

155 154. (Original) The method of claim 152 wherein said model for said alternative hypothesis is:

$$\begin{pmatrix} y_1 \\ y_2 \end{pmatrix} = \begin{pmatrix} \mu_1 \\ \mu_2 \end{pmatrix} + \begin{pmatrix} \beta_1 & \beta_2 \\ \beta_3 & \beta_4 \end{pmatrix} \begin{pmatrix} N_1 \\ N_2 \end{pmatrix} + \begin{pmatrix} \varepsilon_1 \\ \varepsilon_2 \end{pmatrix}$$

wherein

$Q_1$  and  $Q_2$  are categorical random variables indicating the genotype at locus  $Q$  across said plurality of organisms;

$\begin{pmatrix} \varepsilon_1 \\ \varepsilon_2 \end{pmatrix}$  is distributed as a bivariate normal random variable with mean  $\begin{pmatrix} 0 \\ 0 \end{pmatrix}$  and

covariance matrix  $\begin{pmatrix} \sigma_1^2 & \sigma_1\sigma_2 \\ \sigma_2\sigma_1 & \sigma_2^2 \end{pmatrix}$ ;

$\mu_i$  and  $\beta_i$  are model parameters; and one of the conditions (i) through (iv) is valid:

- (i)  $\beta_1 \neq 0$ ,  $\beta_4 \neq 0$ ,  $\beta_2 = 0$ , and  $\beta_3 = 0$ ;
- (ii)  $\beta_1 \neq 0$ ,  $\beta_4 \neq 0$ ,  $\beta_2 \neq 0$ , and  $\beta_3 = 0$ ;
- (iii)  $\beta_1 \neq 0$ ,  $\beta_4 \neq 0$ ,  $\beta_2 = 0$ , and  $\beta_3 \neq 0$ ; and
- (iv)  $\beta_1 \neq 0$ ,  $\beta_4 \neq 0$ ,  $\beta_2 \neq 0$ , and  $\beta_3 \neq 0$ .

155. (Original) The method of claim 151 wherein said comparing comprises:

obtaining a first maximum likelihood estimate for the model for the null hypothesis by maximizing the loglikelihood for the model for the null hypothesis with respect to model parameters;

obtaining a second maximum likelihood estimate for the model for the alternative hypothesis by maximizing the loglikelihood for the model for the alternative hypothesis with respect to model parameters; and

forming a likelihood ratio test statistic between the first maximum likelihood estimate and said second maximum likelihood estimate to determine whether the model for the alternative hypothesis provides for a statistically significant better fit to the data than the model for the null hypothesis.

156. (Original) The method of claim 108 wherein said testing comprises considering a null test for causality having the relationship:

$$P(T_2, Q_* | T_1) = P(T_2 | G)P(Q_* | T_1),$$

wherein

each function  $P$  is a probability density function;

$T_2$  is the variation of the second trait across said plurality of organisms;

$Q_*$  is a genotype random variable for a locus  $Q$  of said one or more loci across said plurality of organisms; and

$T_1$  is the variation of the first trait across said plurality of organisms.

157. (Original) The method of claim 156 wherein said testing comprises comparing said null test for causality, indicating that said first trait  $T_1$  is causal for said second trait  $T_2$ , to an alternative hypothesis, indicating that  $T_2$  and  $Q$  are dependent given  $T_1$ .

158. (Original) The method of claim 157 wherein said testing comprises optimizing the log likelihood ratio of said null hypothesis and said alternative hypothesis using maximum likelihood analysis.

159. (Original) A computer program product for use in conjunction with a computer system, the computer program product comprising a computer readable storage medium and a computer program mechanism embedded therein, the computer program mechanism comprising:

a  $T_1/T_2$  overlap module that comprises instructions for identifying one or more loci in the genome of a species, wherein each locus  $Q$  of said one or more loci is a site of colocalization for (i) a respective quantitative trait locus ( $QTL_1$ ) that is genetically linked to a variation in a first trait  $T_1$  across a plurality of organisms in said species and (ii) a respective quantitative trait locus ( $QTL_2$ ) that is genetically linked to a variation in a second trait  $T_2$  across said plurality of organisms; and

a causality test module that comprises instructions for testing, for one or more locus  $Q$  of said one or more loci, whether (i) a genotype random variable  $Q^*$  of the respective locus  $Q$  across the plurality of organisms and (ii) said variation in the second trait  $T_2$  across the plurality of organisms are correlated conditional on the variation in said first trait  $T_1$  across the plurality of organisms.

160. (Original) A computer system comprising:

a central processing unit;

a memory, coupled to the central processing unit, the memory storing an  $Q_1/Q_2$  overlap module and a causality test module; wherein

the  $T_1/T_2$  overlap module comprises instructions for identifying one or more loci in the genome of a species, wherein each locus  $Q$  of said one or more loci is a site of colocalization for (i) a respective quantitative trait locus ( $QTL_1$ ) that is genetically linked to a variation in the first trait  $T_1$  across a plurality of organisms of said species and (ii) a respective quantitative trait locus ( $QTL_2$ ) that is genetically linked to a variation in the second trait  $T_2$  across said plurality of organisms; and

a causality test module that comprises instructions for testing, for one or more loci  $Q$  in the at least one locus, whether (i) a genotype random variable  $Q^*$  for the respective locus  $Q$  across the plurality of organisms and (ii) said variation in said second trait  $T_2$  across said plurality of organisms are correlated conditional on the variation in the first trait  $T_1$  across said plurality of organisms.

161-210. (Canceled)

211. (Original) A method for determining whether a first trait  $T_1$  is causal for a second trait  $T_2$  in a plurality of organisms of a species, the method comprising:

(A) identifying a locus  $Q$  in the genome of said species that is a site of colocalization for (i) a quantitative trait locus ( $QTL_1$ ) that is genetically linked to a variation in the first trait  $T_1$  across all or a portion of the plurality of organisms and (ii) a quantitative trait locus ( $QTL_2$ ) that is genetically linked to a variation in the second trait  $T_2$  across all or a portion of said plurality of organisms;

(B) quantifying a first coefficient of determination between (i) a genetic variation  $Q^*$  of said locus  $Q$  across all or a portion of said plurality of organisms and (ii) said variation in said first trait  $T_1$  across all or a portion of said plurality of organisms; and

(C) quantifying a second coefficient of determination between (i) said genetic variation  $Q^*$  of said locus  $Q$  across all or a portion of said plurality of organisms and (ii) said variation in said first trait  $T_1$  across all or a portion of said plurality of organisms, after conditioning on said variation in said second trait  $T_2$  across all or a portion of said plurality of organisms, wherein

said first trait  $T_1$  is deemed to be causal for said second trait  $T_2$  when said first coefficient of determination is other than zero and said second coefficient of determination cannot be distinguished from zero.

212. (Currently amended) The method of claim 211 wherein said ~~cellular constituent~~ first trait  $T_1$  is deemed to be causal for said second trait  $T_2$  ~~of interest  $T$~~  when said first coefficient of determination is greater than a predetermined threshold amount.

213. (Original) The method of claim 212 wherein said predetermined threshold amount is 0.03.

214. (Original) The method of claim 212 wherein said predetermined threshold amount is 0.10.

215. (Original) The method of claim 211, wherein said  $QTL_1$  and said  $QTL_2$  are deemed to colocalize at said locus  $Q$  when said  $QTL_1$  and said  $QTL_2$  are within 3 cM of the locus  $Q$ .

216. (Original) The method of claim 211, wherein said  $QTL_1$  and said  $QTL_2$  are deemed to colocalize at said locus  $Q$  when said  $QTL_1$  and said  $QTL_2$  are within 1 cM of the locus  $Q$ .

217. (Original) The method of claim 211 wherein the plurality of organisms is between 100 organisms and 100,000 organisms.

218. (Original) The method of claim 211 wherein the plurality of organisms is less than 500,000 organisms.

219. (Original) The method of claim 211 wherein the plurality of organisms is between 5,000 and 25,000 organisms.

220. (Original) The method of claim 211 wherein said plurality of organisms is human.

221. (Original) The method of claim 211, wherein said first trait  $T_1$  is a complex trait.

222. (Original) The method of claim 221, wherein said complex trait is characterized by an allele that exhibits incomplete penetrance in said species.

223. (Original) The method of claim 221, wherein said complex trait is a disease that is contracted by said at least one organism in said plurality of organisms, and wherein said organism inherits no predisposing allele to said disease.

224. (Original) The method of claim 221, wherein said complex trait arises when one or more of a plurality of different genes in the genome of said species is mutated.

225. (Original) The method of claim 221, wherein said complex trait requires the simultaneous presence of mutations in a plurality of genes in the genome of said species.

226. (Original) The method of claim 221, wherein said complex trait is a phenotype that does not exhibit Mendelian recessive or dominant inheritance attributable to a single gene locus.

227. (Original) The method of claim 221 wherein said complex trait is asthma, ataxia telangiectasia, bipolar disorder, cancer, common late-onset Alzheimer's disease, diabetes, heart disease, hereditary early-onset Alzheimer's disease, hereditary nonpolyposis colon cancer, hypertension, infection, maturity-onset diabetes of the young, mellitus, migraine, nonalcoholic fatty liver, nonalcoholic steatohepatitis, non-insulin-dependent diabetes

mellitus, obesity, polycystic kidney disease, psoriasis, schizophrenia, or xeroderma pigmentosum.

228. (Original) The method of claim 211 wherein said QTL<sub>1</sub> and said QTL<sub>2</sub> are deemed to colocalize at a locus  $Q$  of said one or more loci when said QTL<sub>1</sub> and said QTL<sub>2</sub> are within 40 cM of the locus  $Q$ .

229. (Original) The method of claim 211 wherein said QTL<sub>1</sub> and said QTL<sub>2</sub> are deemed to colocalize at a locus  $Q$  of said one or more loci when said QTL<sub>1</sub> and said QTL<sub>2</sub> are within 10 cM of the locus  $Q$ .

230. (Original) The method of claim 211 wherein said QTL<sub>1</sub> and said QTL<sub>2</sub> are deemed to colocalize at said locus  $Q$  when said QTL<sub>1</sub> and said QTL<sub>2</sub> satisfy a pleiotropy test and wherein failure of the pleiotropy test indicates that the QTL<sub>1</sub> and the QTL<sub>2</sub> are two closely linked QTL and said first trait  $T_1$  is not determined to be causal for said second trait  $T_2$ .

231. (Original) The method of claim 230 wherein said pleiotropy test comprises comparing a model for a null hypothesis, indicating that said QTL<sub>1</sub> and said QTL<sub>2</sub> colocalize as a QTL, to a model for an alternative hypothesis, indicating that said QTL<sub>1</sub> and said QTL<sub>2</sub> are two closely linked QTL.

232. (Original) The method of claim 231 wherein said model for said null hypothesis is:

$$\begin{pmatrix} y_1 \\ y_2 \end{pmatrix} = \begin{pmatrix} \mu_1 \\ \mu_2 \end{pmatrix} + \begin{pmatrix} \beta_1 \\ \beta_2 \end{pmatrix} N + \begin{pmatrix} \varepsilon_1 \\ \varepsilon_2 \end{pmatrix}$$

wherein

$N$  is a categorical random variable indicating the genotype at locus  $Q$  across said plurality of organisms;

$\begin{pmatrix} \varepsilon_1 \\ \varepsilon_2 \end{pmatrix}$  is distributed as a bivariate normal random variable with mean  $\begin{pmatrix} 0 \\ 0 \end{pmatrix}$  and

covariance matrix  $\begin{pmatrix} \sigma_1^2 & \sigma_1\sigma_2 \\ \sigma_2\sigma_1 & \sigma_2^2 \end{pmatrix}$ ; and



$\mu_i$  and  $\beta_i$  are model parameters.

233. (Original) The method of claim 231 wherein said model for said alternative hypothesis is:

$$\begin{pmatrix} y_1 \\ y_2 \end{pmatrix} = \begin{pmatrix} \mu_1 \\ \mu_2 \end{pmatrix} + \begin{pmatrix} \beta_1 & \beta_2 \\ \beta_3 & \beta_4 \end{pmatrix} \begin{pmatrix} N_1 \\ N_2 \end{pmatrix} + \begin{pmatrix} \varepsilon_1 \\ \varepsilon_2 \end{pmatrix}$$

wherein

$N_1$  and  $N_2$  are categorical random variables indicating the genotype at locus  $Q$  across said plurality of organisms;

$\begin{pmatrix} \varepsilon_1 \\ \varepsilon_2 \end{pmatrix}$  is distributed as a bivariate normal random variable with mean  $\begin{pmatrix} 0 \\ 0 \end{pmatrix}$  and

covariance matrix  $\begin{pmatrix} \sigma_1^2 & \sigma_1\sigma_2 \\ \sigma_2\sigma_1 & \sigma_2^2 \end{pmatrix}$ ; and

$\mu_i$  and  $\beta_i$  are model parameters.

234. (Original) The method of claim 231 wherein said model for said alternative hypothesis is:

$$\begin{pmatrix} y_1 \\ y_2 \end{pmatrix} = \begin{pmatrix} \mu_1 \\ \mu_2 \end{pmatrix} + \begin{pmatrix} \beta_1 & \beta_2 \\ \beta_3 & \beta_4 \end{pmatrix} \begin{pmatrix} N_1 \\ N_2 \end{pmatrix} + \begin{pmatrix} \varepsilon_1 \\ \varepsilon_2 \end{pmatrix}$$

wherein

$Q_1$  and  $Q_2$  are categorical random variables indicating the genotype at locus  $Q$  across said plurality of organisms;

$\begin{pmatrix} \varepsilon_1 \\ \varepsilon_2 \end{pmatrix}$  is distributed as a bivariate normal random variable with mean  $\begin{pmatrix} 0 \\ 0 \end{pmatrix}$  and

covariance matrix  $\begin{pmatrix} \sigma_1^2 & \sigma_1\sigma_2 \\ \sigma_2\sigma_1 & \sigma_2^2 \end{pmatrix}$ ;

$\mu_i$  and  $\beta_i$  are model parameters; and one of the conditions (i) through (iv) is valid:

(i)  $\beta_1 \neq 0$ ,  $\beta_4 \neq 0$ ,  $\beta_2 = 0$ , and  $\beta_3 = 0$ ;

(ii)  $\beta_1 \neq 0$ ,  $\beta_4 \neq 0$ ,  $\beta_2 \neq 0$ , and  $\beta_3 = 0$ ;

(iii)  $\beta_1 \neq 0$ ,  $\beta_4 \neq 0$ ,  $\beta_2 = 0$ , and  $\beta_3 \neq 0$ ; and

(iv)  $\beta_1 \neq 0$ ,  $\beta_4 \neq 0$ ,  $\beta_2 \neq 0$ , and  $\beta_3 \neq 0$ .

235. (Original) The method of claim 231 wherein said comparing comprises:

obtaining a first maximum likelihood estimate for the model for the null hypothesis by maximizing the loglikelihood for the model for the null hypothesis with respect to model parameters;

obtaining a second maximum likelihood estimate for the model for the alternative hypothesis by maximizing the loglikelihood for the model for the alternative hypothesis with respect to model parameters; and

forming a likelihood ratio test statistic between the first maximum likelihood estimate and said second maximum likelihood estimate to determine whether the model for the alternative hypothesis provides for a statistically significant better fit to the data than the model for the null hypothesis.

236. (Currently amended) A method for identifying a quantitative trait locus for a trait that is exhibited by a plurality of organisms in a population, comprising:

(a) dividing said population into a plurality of sub-populations using a classification scheme that classifies each organism in said population into at least one of said subpopulations, wherein said classification scheme is derived from a plurality of cellular constituent measurements for each of a plurality of respective cellular constituents that are obtained from each said organism and ~~wherein~~ wherein said classification scheme uses a classifier constructed using boosting or adaptive boosting; and

(b) for at least one sub-population in said plurality of sub-populations, performing quantitative genetic analysis on said sub-population in order to identify said quantitative trait locus for said trait.

237-296. (Canceled)

297. (New) A computer program product for use in conjunction with a computer system, the computer program product comprising a computer readable storage medium and a computer program mechanism embedded therein, the computer program mechanism for determining

whether a first trait  $T_1$  is causal for a second trait  $T_2$  in a plurality of organisms of a species, the computer program mechanism comprising:

(A) instructions for identifying a locus  $Q$  in the genome of said species that is a site of colocalization for (i) a quantitative trait locus ( $QTL_1$ ) that is genetically linked to a variation in the first trait  $T_1$  across all or a portion of the plurality of organisms and (ii) a quantitative trait locus ( $QTL_2$ ) that is genetically linked to a variation in the second trait  $T_2$  across all or a portion of said plurality of organisms;

(B) instructions for quantifying a first coefficient of determination between (i) a genetic variation  $Q^*$  of said locus  $Q$  across all or a portion of said plurality of organisms and (ii) said variation in said first trait  $T_1$  across all or a portion of said plurality of organisms; and

(C) instructions for quantifying a second coefficient of determination between (i) said genetic variation  $Q^*$  of said locus  $Q$  across all or a portion of said plurality of organisms and (ii) said variation in said first trait  $T_1$  across all or a portion of said plurality of organisms, after conditioning on said variation in said second trait  $T_2$  across all or a portion of said plurality of organisms, wherein

said first trait  $T_1$  is deemed to be causal for said second trait  $T_2$  when said first coefficient of determination is other than zero and said second coefficient of determination cannot be distinguished from zero.

298. (New) A computer system comprising:

a central processing unit;

a memory, coupled to the central processing unit, the memory comprising:

(A) instructions for identifying a locus  $Q$  in the genome of said species that is a site of colocalization for (i) a quantitative trait locus ( $QTL_1$ ) that is genetically linked to a variation in the first trait  $T_1$  across all or a portion of the plurality of organisms and (ii) a quantitative trait locus ( $QTL_2$ ) that is genetically linked to a variation in the second trait  $T_2$  across all or a portion of said plurality of organisms;

(B) instructions for quantifying a first coefficient of determination between (i) a genetic variation  $Q^*$  of said locus  $Q$  across all or a portion of said plurality of organisms and (ii) said variation in said first trait  $T_1$  across all or a portion of said plurality of organisms; and

(C) instructions for quantifying a second coefficient of determination between (i) said genetic variation  $Q^*$  of said locus  $Q$  across all or a portion of said plurality of

organisms and (ii) said variation in said first trait  $T_1$  across all or a portion of said plurality of organisms, after conditioning on said variation in said second trait  $T_2$  across all or a portion of said plurality of organisms, wherein

said first trait  $T_1$  is deemed to be causal for said second trait  $T_2$  when said first coefficient of determination is other than zero and said second coefficient of determination cannot be distinguished from zero.

299. (New) The method of claim 108, wherein said second trait  $T_2$  is a complex trait.

300. (New) The method of claim 299, wherein said complex trait is characterized by an allele that exhibits incomplete penetrance in said species.

301. (New) The method of claim 299, wherein said complex trait is a disease that is contracted by an organism in said plurality of organisms, and wherein said organism inherits no predisposing allele to said disease.

302. (New) The method of claim 299, wherein said complex trait arises when any of a plurality of different genes in the genome of said species are mutated.

303. (New) The method of claim 299, wherein said complex trait requires the simultaneous presence of mutations in a plurality of genes in the genome of said species.

304. (New) The method of claim 299, wherein said complex trait is associated with a high frequency of disease-causing alleles in said species.

305. (New) The method of claim 299, wherein said complex trait is a phenotype that does not exhibit Mendelian recessive or dominant inheritance attributable to a single gene locus.

306. (New) The method of claim 299, wherein said complex trait is asthma, ataxia telangiectasia, bipolar disorder, cancer, common late-onset Alzheimer's disease, diabetes, heart disease, hereditary early-onset Alzheimer's disease, hereditary nonpolyposis colon cancer, hypertension, infection, maturity-onset diabetes of the young, mellitus, migraine,

nonalcoholic fatty liver, nonalcoholic steatohepatitis, non-insulin-dependent diabetes mellitus, obesity, polycystic kidney disease, psoriasis, schizophrenia, or xeroderma pigmentosum.

307. (New) The method of claim 109, wherein said first trait is abundance levels of a first cellular constituent and each quantitative measurement of said first trait is an abundance level of said first cellular constituent in an organism in said plurality of organisms.

308. (New) The method of claim 211, wherein said second trait  $T_2$  is a complex trait.

309. (New) The method of claim 308, wherein said complex trait is characterized by an allele that exhibits incomplete penetrance in said species.

310. (New) The method of claim 308, wherein said complex trait is a disease that is contracted by an organism in said plurality of organisms, and wherein said organism inherits no predisposing allele to said disease.

311. (New) The method of claim 308, wherein said complex trait arises when any of a plurality of different genes in the genome of said species are mutated.

312. (New) The method of claim 308, wherein said complex trait requires the simultaneous presence of mutations in a plurality of genes in the genome of said species.

313. (New) The method of claim 308, wherein said complex trait is associated with a high frequency of disease-causing alleles in said species.

314. (New) The method of claim 308, wherein said complex trait is a phenotype that does not exhibit Mendelian recessive or dominant inheritance attributable to a single gene locus.

315. (New) The method of claim 308, wherein said complex trait is asthma, ataxia telangiectasia, bipolar disorder, cancer, common late-onset Alzheimer's disease, diabetes, heart disease, hereditary early-onset Alzheimer's disease, hereditary nonpolyposis colon

cancer, hypertension, infection, maturity-onset diabetes of the young, mellitus, migraine, nonalcoholic fatty liver, nonalcoholic steatohepatitis, non-insulin-dependent diabetes mellitus, obesity, polycystic kidney disease, psoriasis, schizophrenia, or xeroderma pigmentosum.

316. (New) The method of claim 212, wherein said first trait is abundance levels of a first cellular constituent and each quantitative measurement of said first trait is an abundance level of said first cellular constituent in an organism in said plurality of organisms.

317. (New) The method of claim 108, wherein

the first trait  $T_1$  is an abundance level of a cellular constituent;

each said respective quantitative trait locus ( $QTL_1$ ) that is genetically linked to a variation in the first trait  $T_1$  across the plurality of organisms is a respective abundance quantitative trait locus (eQTL) that is genetically linked to a variation in abundance levels of the cellular constituent across the plurality of organisms;

the second trait  $T_2$  is a trait of interest  $T$  exhibited by one or more organisms in the plurality of organisms; and

each said respective quantitative trait locus ( $QTL_2$ ) that is genetically linked to a variation in the second trait  $T_2$  is a respective clinical quantitative trait locus (cQTL) that is genetically linked to a variation in the trait of interest  $T$  across the plurality of organisms.

318. (New) The method of claim 211, wherein

the first trait  $T_1$  is an abundance level of a cellular constituent;

each said respective quantitative trait locus ( $QTL_1$ ) that is genetically linked to a variation in the first trait  $T_1$  across the plurality of organisms is a respective abundance quantitative trait locus (eQTL) that is genetically linked to a variation in abundance levels of the cellular constituent across the plurality of organisms;

the second trait  $T_2$  is a trait of interest  $T$  exhibited by one or more organisms in the plurality of organisms; and

each said respective quantitative trait locus ( $QTL_2$ ) that is genetically linked to a variation in the second trait  $T_2$  is a respective clinical quantitative trait locus (cQTL) that is genetically linked to a variation in the trait of interest  $T$  across the plurality of organisms.